

Project MULTRA, Subproject 135

Date Initiated: 15 September 1961

Date Expires: 1 October 1962

Funds-current year: \$24,870.00

Purpose: To provide for investigations involving clinical testing of volunteers in programs of TED interest. The project will provide a facility and a team of medical supervisors as well as a large stable reservoir of volunteers.

Status: Project is approved but has not been activated. Activation ... will be attempted by I September 1961.



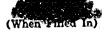
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Cost Account \_\_ 3/35- /390 - 3902

Object Class .

Date	Remarks and References	Obligations Incurred	Obligations Liquidated	Unliquidated Balance
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16 June 1961

MEMORANDUM FOR: COMPTROLLER	
ATTENTION : Finance Division	
SUBJECT : MKULTRA, Subproject 135	
	•
Under the authority granted in the memorandum dated 16 June	
1961 from the DCI to the DD/A, and the extension of this authority in	
subsequent memoranda, Subproject 135 has been approved and \$25, 86	18.00
of the over-all MKULTRA project funds has been obligated to cover the	he
subproject expenses. This obligation of funds should be charged to	
Allotment 2125-1390-3902.	
TSD/Research Branch	
APPROVED FOR	
OBLIGATION OF FUNDS:	
Original signed by Certify that funds are ayanable Grugation affective no. 132	and the second s
Sharet to augment no. 2125-12	190.3902
AUTHORIZING OFFICER	
Research Director	
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Date:	e de la companya de
Distribution;	
Orig & 2 - Addresse e	
1 - TSD/OC	
TSD/FASS	
2 - TSD/RB	



13 YILY 1961

KENORANDUM FOR: CHIEF, PINANCE DIVISION

VIA !

TSD/Budget Officer

SUBJECT

: MOULTRA, Subproject 135, Invoice fl Allotment 2125-1390-3902.

1. Invoice No. 1 covering the above subproject is attached. Payment should be made as follows:

back in the angunt of 124 200 .00 .4m payable In the amount of 1000 and mayer bayable to

2. Please forward the checks to Chief, TSD/Research Branch, through TSD/Budget Officer, no later than 24 July 1961.

3. This is a final invoice. However, since it is anticipated that additional funds vill be obligated for this project, the files should not be closed.

TSD/Research Branch \*

Attachments:

Invoice & Certifications

2 - TSD/RB

CERTIFY THAT PUNDS ARE AVAILABLE

CRUGATION REFERENCE No.

CHAPCE TO ALL DE MENT HO. 2/25 1280 - 3902-

Distributions

Orig & 2 - Addresses

1 - TSD/FASS

AUTHORIZING OFFICER

Rirhl



INVOICE

For Services

\$25,868.00



### CERTIFICATIONS

(1) It is hereby certified that this is Invoice 1 applying to sub-project No. 135 of MKULTRA, that performance is satisfactory, that services are being accomplished in accordance with mutual agreements, that a detailed agenda of the payments and receipts is on file in TSD/RB, that this bill is just and correct and that payment thereof has not yet been made.

Chief, TSD/Research Branch

		And the second s	·
(s)	It is	hereby certified that this invoice	applies to SubProject 135

of MCULTRA which was duly approved, and that the project is being continuous in accordance with the memorandum of 13 April 1953 from the DCI to the DD/A, and the extension of this authority in subsequent memoranda.

Research Director

Date

Data.

(When Filled In)

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11 June 1964

MEMORATEDISM FOR 1 THE RELICED

SUBJECT

: MAILTRA Subprojects 70, 91, 94 and 135

Since the completion of the audit of Project MULTRA for the period 1 August 1060 through 31 Jenuary 1963, accountings have been received from the should be should be should be accounted for he grantees at the end of the audit period.

Subproject 70 incurred direct costs of \$2,518.34 and \$27,872.16 transferred from Subproject 94, for a total of \$30,390.50. Subproject 94 incurred expenses of \$71,324.87, but transferred \$4,165.71 to Subproject 91 and \$27,872.16 to Subproject 70, leaving a not cost of \$39,287.00. It was decided to divert the finds for Subproject 135 to Subproject 91, therefore these costs were included in the accounting from Subproject 91.

Innumber as: 1) all four of the subprojects had the some basic research goal in mind, 2) the great length of time that has elapsed since the grants were made, and 3) the difficulty in getting accountings of any sort from the grantes, it is recommended that we facilitate this matter by accounting for the balances as above in the Confidential Funds Division 760 account for these subprojects.

The above has been concurred in by

the Audit

Chief Scientify was



# Pharmacological Screening and Testing (Proj.500w) December 1, 1961 to July 31, 1962

February 1, 1961 to January 31, 1962 Less, previously reported	\$\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	\$8,000.00
February 1, 1962 to July 31, 1962		24,000.00-135
<u>Total</u>		\$32,000.00 (N SUB 91)
Accounted for, as follows:		\$28,131.14
Salaries		1,400.08
Payroll Taxes	<del></del>	355.68
Employee Benefits Laboratory Supplies and Expense Travel		3,541.48 4 569.61
Other Direct Costs		2,569.17
Indirect Costs		36,372.26
Total Costs	<u> </u>	72,939.42
Less, Previously reported Total Accounted For		39,894.36 \$33,045.06





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# **CHARLES**

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16 August 1961



Gentlemen:

We are pleased to be able to transmit to you the following

funds!



These funds represent a contribution for the use of your directors in carrying out the very worthwhile research goals of your obligations.

Yours truly.

Attachments (2)



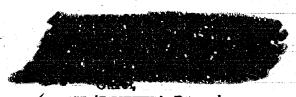
D R A F T 31 May 1962

MEMO	ANITHE	A TOP.	THE RE	CORD
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SUBJECT

Project MKULTRA, Subproject No. 135, Change of Facility in which Work is Performed

- 1. The original purpose of this subproject was to evaluate a number of candidate chemical agents in facilities and under the direction of individuals who, it later developed, became unavailable for this purpose due to unforeseen circumstances.
- 2. In the meantime, the funds for this subproject were transferred to the cutout in anticipation of the commencement of the work.
- 3. In view of the security difficulties involved in the actual return of funds from the cutout and also since another subproject (#91) was being served by the same cutout, and furthermore, since subproject #91 was capable of providing its services on these chemical agents of interest and during the same fiscal year, it has been decided to divert the work intended for subproject #135 to the facilities of subproject #91. No significant change in funding or variation in fiscal year is concerned in this matter.



TSD/Research Branch

CONCUR:



Distribution:

Original only

Date:



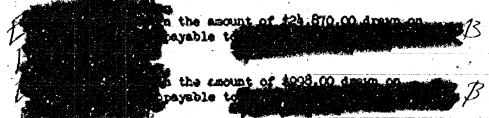


12 July 1961

MEMORANDUM	OR: CHIKP, PINANCE DIVISION
VIA :	1 TSD/Budget Officer
SUBJECT	Allotment 2125-1390-3902.

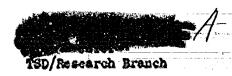
1. Invoice No. 1 covering the above subproject is attached.

Payment should be made as follows:



2. Please forward the checks to Chief, TSD/Research Branch, through TSD/Budget Officer, no later than 24 July 1961.

3. This is a final invoice. However, since it is anticipated that additional funds will be obligated for this project, the files should not be closed.



Attachments:	 
Invoice & Certification	
Distribution	

1 - TSD/FASS

Orig & 2 - Addresses







\$25,868.00 For Services CERTIFICATIONS (1) It is hereby certified that this is Invoice 1 applying to sub-project No. 135 of MOULTRA, that performance is satisfactory, that services are being accomplished in accordance with mutual agreements, that a detailed agenda of the payments and receipts is on file in TSD/RB, that this bill is just and correct and that payment thereof has not yet been made. Chief, TSD/Research Branch Date: (2) It is hereby certified that this invoice applies to SubProject 135 of MKULTRA which was duly approved, and that the project is being carried out in accordance with the memorandum of 13 April 1953 from the DCI to the DD/A, and the extension of this authority in subsequent memoranda. Research Director

Date:



16 June 1961

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MEMORANDUM	FOR CO	MPTROLLES			
ATTENTION	ıFin	ance Division	3		
SUBJECT	; MK	ULTRA, Sub	project 135		
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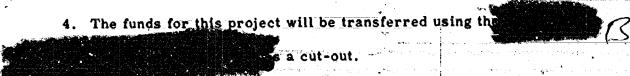
16 June 1961

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SUBJECT

MKULTRA, Subproject 135

- 1. The purpose of this subproject is to provide funds to carry out a research program involving utilization of human volunteer subjects.
- 2. This program is designed to exploit clinical and laboratory
  facilities and professional knowledge in providing answers to specific
  questions of direct interest to the Agency. The nature of the investigations to be carried out encompasses pharmacological and clinical
  testing in human volunteers of chemicals, biochemicals, and biologically
  active materials under controlled conditions. For example, human
  patch testing or human blood group determinations will be mounted upon
  setting up a suitably designed experiment.



5. The cost of this program for a period of one year beginning 1 July 1961 will not exceed \$24,870.00. To this sum must be added \$998.00 representing a 4% service charge to the

who will act as a cut-out, making the total amount \$25,868.00.

Charges should be made against allotment #2125-1390-3902.

- pal investigator who administers the grant to submit to them a summary accounting upon an annual basis of all monies received from the Also they will require that any unexpended funds be returned as soon as possible upon the close of the grant year.
- 7. The requirement for a six months' informal accounting on the part of the principal investigator is waived.
- 8. It is not anticipated that any permanent equipment will be required for this program.
- 9. It has been agreed that documentation and accounting for travel expenses which are reimbursable shall conform to accepted practices.
- 10. It is agreed that technical reports reflecting the progress of the research program will be accepted at mutually accepted intervals.
- TOP SECRET Agency Security Agreement. The project will be unclassified after it reaches the

B





Chief, TSD/Research Branch

APPROVED FOR OBLIGATION OF FUNDS:



Research Director

Date: 12101

Attachment:

Proposal & Budget

Distribution:
Original only

# **PROPOSAL**

# OBJECTIVE :

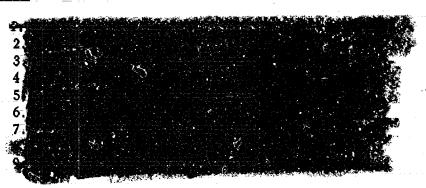
To establish a professional organization for the primary purpose of servicing and conducting complete clinical evaluations and testings upon demand.

In addition to prescribed activities, to maintain itself on a standby basis and to be able to mount clinical projects on an ad hoc basis.

PROCEDURE:

see attached sample protocol for patch testing.

# CLINICAL FACILITIES:



# LABORATORY FACILITIES:



PROPESSIONAL STAFF:

# BUDGET: (12 months period)

Principal investigator	10,000.00
Professional staff	6,200.00
Secretarial	1,800.00
Expendables, chemicals, supplies	1,600.00
Clinical subjects, fees	2,500.00
Travel, maintenance	2,300.00
Insurance, malpractice & liability	470.00
Total	24,870.00

no overhead or fee

### PROTOCOL

# **OBJECTIVE:**

To determine whether an unknown compound, either chemical or biological, is capable of eliciting a reaction when applied topically to the skin, either locally or systemically, or both.

# **BACKGROUND:**

If the skin is able to deal with the irritant compound, the reaction is physiological or normal. However, if the action of the irritant is excessive, the reaction is pathological or abnormal and irritation results. Irritation is an extreme reaction of tissues to an insult or injury and is best characterized as incipient inflammation. The inflammatory process may vary from a barely perceptible hyperemia, to edema formation and vesiculation and finally to a suppurative process.

A primary irritant may be defined as a substance producing an injury on first contact. The resultant injury will depend on the nature of the irritant, concentration of the irritant, and the total elapsed time of exposure. Primary irritation of the skin is usually measured experimentally by the patch test technique on abraded or intact skin of the experimental animal, for example, rabbits, guinea pigs or man.

Exposure of man is predicated on small animal studies used to screen all compounds to be tested to obviate exposing man to severely sensitizing substances. Granted, in the case of compounds which may be only mildly sensitizing or in which small animal studies are unable to furnish unequivocal results, man may appear to be the only feasible final test subject.

Animal experimentation data should include acute, subacute and chronic dermal toxicity experimentation that will allow some logical assessment of the local and systemic toxicity of substances to be applied topically to man in the proper amounts, concentration and frequency of application that may be tolerated.

### PROCEDURE:

per compound tested. Prior to experimental testing, maximum efforts must be made to minimize the risk of incurring unforseeable sensitivities and untoward responses by establishing a base line profile that will encompass the following:

- 1. Complete medical histories with emphasis on known sensitivities and drug idiosyncracles.
- 2. Preliminary physicial examination
- 3. Hematological evaluation ( complete blood count )
- 4. Urinalysis ( microscopic )
- 5. Radiological screening (chest and long bone)

Acute dermal toxicity. (Single exposure or primary irritation)

This will utilize the patch test technique on intact skin on a minimum of five subjects per test compound.

Methodology will consist of introducing under a one inch patch 0.5 ml (liquid) or 0.5 gm (solid or semi-solid) of test substance and securing the patch in place with non-sensitizing adhesive.

After 24 hours, the patch is removed and readings made on the resultant reactions evaluated on the basis of the following scores:

Evaluation of Skin Reactions

1.	Brythema and Eschar Formation	
	No erythema	0
	Very slight erythema (barely perceptible)	1
	Well defined erythema	2
:	Moderate to severe erythema	3 11 71 11 11 11 11
:	Severe erythema (beet redness) to	4
	Slight eschar formation (injuries in depth)	. <u>4</u>
-	Total possible erythema score	4.1.1.
Ž.	- Edema Formation	
	No edema	0
•	Very slight edema (barely perceptible)	
		_2
:	Moderate edema (raised approx 1 mm)	<b>3</b>
	Severe edema (raised more than 1 mm and	
	extending beyond area of exposure)	4
	Total possible edema score	4

Readings are again made after 72 hours of exposure. The final score represents the average of 24-72 hour readings.

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